

PTD

IMPROVED CATALYTIC COMPOSITION AND ITS APPLICATION TO OLEFIN OLIGOMERIZATION

Field of the Invention

The present invention relates to a novel catalytic composition for use in oligomerization processes, in particular for monoolefin dimerization.

It also relates to a process for oligomerization, in particular dimerization, of monoolefins, using such a catalytic composition.

Description of the Prior Art

The preparation of catalysts for dimerization or co-dimerization of monoolefins such as ethylene, propylene, butenes or pentenes is known. Examples of such catalysts which can be cited are the products of the interaction of π -allyl nickel phosphine halides with Lewis acids (French patent FR-B-1 410 430), the products of the interaction of nickel phosphine halides with Lewis acids (United States patent US-A-3 485 881) and the products of the interaction of certain nickel carboxylates with hydrocarbylaluminium halides (US-A-3 321 546).

Almost all of those catalysts use a ligand such as an organic phosphorus compound and it is preferable to have phosphorus-free oligomerization catalysts available. To this end, it is possible to use zerovalent nickel compounds but they are of little practical use because of their instability. It is also possible to use catalysts in which the nickel is deposited on a mineral support comprising acid sites, such as silica, alumina or silica-aluminas. However, in that case the catalysts are solid, in contrast to the desired liquid catalysts.

US-A-4 283 305 teaches that an association of a divalent nickel compound, a hydrocarbyl-aluminium halide with formula AlR_mX_{3-m} where R is a hydrocarbyl radical containing 1 to 12 carbon atoms, X is a chlorine or bromine atom, and m is a number from 1 to 2, and a compound with a Bronsted acid nature, leads to a catalytic composition that is more active than previously, and also less sensitive to the trace impurities which are routinely found in industrial olefinic feeds.

Summary of the Invention

It has now, unexpectedly, been found that for a catalytic composition obtained by bringing a divalent nickel compound into contact with an organic Bronsted acid and a hydrocarbylaluminium

dihalide, pre-conditioning the catalytic composition in a solvent before using it for oligomerization can still further increase the activity for olefin oligomerization. The use of a hydrocarbylaluminium dihalide enriched with an aluminium trihalide can still further increase the activity of the catalytic composition.

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Detailed Description of the Invention

More precisely, said improved catalytic composition comprises the product resulting from bringing the following three constituents into contact in any order:

- a) at least one divalent nickel compound;
- b) at least one hydrocarbylaluminium dihalide with formula AlRX_2 , where R is a hydrocarbyl radical containing 1 to 12 carbon atoms such as alkyl, aryl, aralkyl or cycloalkyl, and X is a chlorine or bromine atom; and
- c) at least one organic Bronsted acid;

the mixture obtained being pre-conditioned in a solvent, at a controlled temperature and for a pre-set period, prior to its use.

The divalent nickel compound can be any compound soluble in a proportion of more than 1 g per litre in a hydrocarbon medium, more particular in the reactants and the reaction medium. Preferably, nickel carboxylates are used with general formula $(\text{R}_1\text{COO})_2\text{Ni}$, where R_1 is a hydrocarbyl radical, for example alkyl, cycloalkyl, alkenyl, aryl, aralkyl or alkaryl, containing up to 20 carbon atoms, preferably a hydrocarbyl residue containing 5 to 20 carbon atoms. Radical R_1 can be substituted by one or more halogen atoms, hydroxyl groups, ketone, nitro, cyano or other groups that do not interfere with the reaction. The two radicals R_1 can also constitute an alkylene residue containing 6 to 18 carbon atoms. Non limiting examples of nickel compounds are the following divalent nickel salts: octoate, 2-ethylhexanoate, decanoate, stearate, oleate, salicylate and hydroxy-decanoate. Preferably, nickel 2-ethylhexanoate is used.

The Bronsted acid compound has formula HY , where Y is an organic anion, for example carboxylic, sulphonic or phenolic. Preferably, acids with a pK_a of a maximum of 3 at 20°C are used, more particularly those which are soluble in the nickel compound or in its solution in a hydrocarbon or another suitable solvent. One preferred class of acids comprises the group formed by

halogenocarboxylic compounds with formula R_2COOH where R_2 is a halogenated alkyl radical, in particular those which contain at least one halogen atom alpha to the $-COOH$ group, with a total of 2 to 10 carbon atoms. Preferably, a halogenoacetic acid with formula $CX_pH_{3-p}COOH$ is used where X is fluorine, chlorine, bromine or iodine, and p is a whole number from 1 to 3. Examples which 5 can be cited are trifluoroacetic, difluoroacetic, fluoroacetic, trichloroacetic, dichloroacetic and chloroacetic acids. These examples are not limiting and it is also possible to use arylsulphonic acids, alkylsulphonic acids, fluoroalkylsulphonic acids, picric acid and nitroacetic acid. Preferably, trifluoroacetic acid is used.

10 The three constituents of the catalytic formula can be mixed in any order. However, it is preferable to first mixture the nickel compound with the organic Bronsted acid then to introduce the aluminium compound. The mole ratio of the hydrocarbylaluminium dihalide to the nickel compound, expressed as the Al/Ni ratio, is 2/1 to 50/1, preferably 2/1 to 20/1. The mole ratio of the Bronsted acid to the nickel compound is 0.25/1 to 10/1, preferably 0.25/1 to 5/1.

15 Pre-conditioning the catalytic composition consists of mixing the three constituents in a hydrocarbon solvent, for example in an alkane or in an aromatic hydrocarbon, or in a halogenated hydrocarbon or, as is preferred, in a mixture with a composition analogous to that of the mixtures obtained in the dimerization or oligomerization reaction itself. Thus for a catalytic composition intended for propylene dimerization, the pre-conditioning solvent can principally be constituted by isohexenes.

20 The mixture is generally produced by stirring in an inert atmosphere, for example in nitrogen or argon, at a controlled temperature of 0°C to 80°C, preferably 10°C to 60°C, for a period of 1 minute to 5 hours, preferably 5 minutes to 1 hour. The solution obtained is then transferred into the oligomerization reactor under an inert atmosphere.

25 In a preferred implementation, in the catalytic composition of the invention, the hydrocarbylaluminium dihalide can be enriched with an aluminium trihalide, the mixture of the two compounds then having formula AlR_nX_{3-n} , where R and X are as defined above and n is a number from 0 to 1 (limits excluded).

The hydrocarbylaluminium dihalide compounds enriched in aluminium trihalide are obtained by mixing a hydrocarbylaluminium dihalide with formula AlRX_2 where R is a hydrocarbyl radical containing 1 to 12 carbon atoms, such as alkyl, aryl, aralkyl, alkaryl or cycloalkyl and X is a chlorine atom or a bromine atom, with an aluminium trihalide AlX_3 . Non limiting examples of such compounds which can be cited are: dichloroethylaluminium enriched with aluminium trichloride, the mixture having formula $\text{AlEt}_{0.9}\text{Cl}_{2.1}$, for example; dichloroisobutylaluminium enriched with aluminium trichloride, the mixture having formula $\text{AlBu}_{0.9}\text{Cl}_{2.1}$, for example, and dibromoethylaluminium enriched with aluminium tribromide, the mixture having formula $\text{AlEt}_{0.9}\text{Br}_{2.1}$, for example.

In this case as well, the three constituents of the catalytic formula can be mixed in any order. It is also preferable to first mix the nickel compound with the organic Bronsted acid, then to introduce the aluminium compound. In this case, it is = the mole ratio between the hydrocarbylaluminium dihalide enriched with aluminium trihalide and the nickel compound, expressed as the Al/Ni ratio, which is 2/1 to 50/1, preferably 2/1 to 20/1. As indicated above, the mole ratio of the Bronsted acid to the nickel compound is still 0.25/1 to 10/1, preferably 0.25/1 to 5/1.

The invention also relates to a process for oligomerization, in particular dimerization, of monoolefins in the presence of the catalytic system defined above.

Examples of monoolefins that can be dimerized or oligomerized are ethylene, propylene, butenes, pentenes and hexenes, used pure or as a mixture, contained in cuts from refining or from 20 chemistry. These olefins can also be co-oligomerized between themselves.

The process can be carried out in a reactor with one or more reaction stages in series, the olefinic feed and/or the previously pre-conditioned catalytic composition being introduced continuously, either to the first stage, or to the first and to any other stage.

The process is generally carried out at a temperature of -20°C to $+80^\circ\text{C}$, under pressure 25 conditions such that at least the major portion of the reactants are maintained in the liquid phase or in the condensed phase.

At the reactor outlet, the catalyst can be deactivated, for example by injecting ammonia and/or an aqueous sodium hydroxide solution and/or an aqueous sulphuric acid solution. The

unconverted olefins and alkanes which may be present in the feed are then separated from the oligomers by distillation.

The products obtained by the process of the invention can be used, for example, as constituents for automobile fuels and/or as feeds for a hydroformylation process for synthesising 5 aldehydes and alcohols.

The entire disclosure of all applications, patents and publications, cited above and below, and of corresponding French application 99/06749, filed on May 27, 1999 are hereby incorporated by reference.

The following examples illustrate the invention without in any way limiting its scope.

10 **EXAMPLE 1 (comparative)**

Preparation of catalyst:

In this example of the prior art, the catalyst was prepared in situ in the autoclave where oligomerization took place, without pre-conditioning, and used dichloroethylaluminium as the aluminium compound.

15 Use in oligomerization:

A solution of 0.043 g of nickel 2-ethylhexanoate containing 13% by weight of nickel in 40 ml of isohexene solvent was introduced into a 250 ml stainless steel autoclave provided with stirring and wherein the temperature could be regulated by circulating water in an external envelope, followed by 10 g of a mother liquor prepared from 0.11 g of trifluoroacetic acid made up to 100 g 20 with isohexene solvent, introduced with stirring. Finally, a solution of 0.18 g of dichloroethylaluminium in 50 ml of isohexene solvent was injected. This corresponded to 1.41 mmole of aluminium and an Al/Ni mole ratio of 15/1.

Subsequently, 10 g of isohexenes was added to the autoclave to make up the solvent, followed by 10.8 g of liquid propylene via pressure lock. The temperature was rapidly raised to 25 40°C. After 15 minutes of reaction, the conversion of propylene into a mixture of dimers, trimers and tetramers containing 80% by weight of dimers was 67%.

EXAMPLE 2 (in accordance with the invention)

Preparation of catalyst:

0.043 g of nickel 2-ethylhexanoate containing 13% by weight of nickel was introduced into a 250 ml glass flask provided with a magnetic stirrer, then the flask was carefully purged and placed under an argon atmosphere. A transfer needle was used to introduce 40 ml of a fraction of isohexenes distilled under argon and dried over 3A molecular sieve, which was then used as the solvent. 5 Stirring dissolved the nickel salt. 10 g of a mother liquor prepared from 0.11 g of trifluoroacetic acid made up to 100 g with the isohexene solvent was then injected. This was all placed, with continued stirring, in a thermostatic bath regulated to 30°C.

10 A further flask purged with argon was used to prepare a solution of 0.18 g of dichloroethylaluminium in 50 ml of isohexene solvent. The solution obtained was slowly added to the nickel solution prepared above using a transfer needle. This corresponded to 1.41 mmole of total aluminium and to a Al/Ni mole ratio of 15/1. This was all pre-conditioned at 30°C for 30 minutes with stirring.

Use in oligomerization:

15 The pre-conditioned catalytic solution was transferred under argon to an autoclave as described in Example 1. 10 g of isohexenes was then introduced into the autoclave to make up the solvent, followed by 10.8 g of liquid propylene using a pressure lock. The temperature was rapidly raised to 40°C. The reaction was followed by periodically removing samples for gas chromatographic analysis. After 15 minutes of reaction, the conversion of propylene into a mixture of dimers, trimers and tetramers analogous to that of Example 1 was 86%.

20 **EXAMPLE 3 (in accordance with the invention)**

Preparation of catalyst:

0.043 g of nickel 2-ethylhexanoate containing 13% by weight of nickel was introduced into a 250 ml glass flask provided with a magnetic stirrer, then the flask was carefully purged and placed under an argon atmosphere. A transfer needle was used to introduce 40 ml of a fraction of isohexenes distilled under argon and dried over 3A molecular sieve, which was then used as the solvent. 25 Stirring dissolved the nickel salt. 10 g of a mother liquor prepared from 0.11 g of trifluoroacetic acid made up to 100 g with the isohexene solvent was then injected. This was all placed, with continued stirring, in a thermostatic bath regulated to 30°C.

50 ml of isohexene solvent was introduced into a further flask purged with argon, followed by 0.165 g of dichloroethylaluminium and finally 0.015 g of aluminium trichloride, corresponding to a compound with formula $\text{AlEt}_{0.92}\text{Cl}_{2.08}$. The solution obtained was slowly added to the nickel solution prepared above using a transfer needle, whereupon the colour changed from green to yellow. This corresponded to 1.41 mmole of total aluminium and to an Al/Ni mole ratio of 15/1. This was all pre-conditioned at 30°C for 30 minutes with stirring.

Use in oligomerization:

10 The pre-conditioned catalytic solution was transferred under argon to an autoclave as described in Example 1. 10 g of isohexenes was then introduced into the autoclave to make up the solvent, then 10.8 g of liquid propylene using a pressure lock. The temperature was rapidly raised to 40°C. The reaction was followed by periodically removing samples for gas chromatographic analysis. After 15 minutes of reaction, the conversion of propylene into a mixture of dimers, trimers and tetramers analogous to that of Example 1 was 89%.

15 The preceding examples can be repeated with similar success by substituting the generically or specifically described reactants and/or operating conditions of this invention for those used in the preceding examples.

20 From the foregoing description, one skilled in the art can easily ascertain the essential characteristics of this invention and, without departing from the spirit and scope thereof, can make various changes and modifications of the invention to adapt it to various usages and conditions.